AMENDMENTS TO THE CLAIMS

- 1-2. (Canceled)
- 3. (Previously presented) The composition of Claim 34, wherein the therapeutic, diagnostic, or prophylactic agent is a protein, peptide, nucleotide, oligonucleotide, saccharide, polysaccharide, organic molecule, or combination thereof.
- 4. (Currently amended) The composition of Claim [[33]] <u>36</u>, wherein the hydrophobic component is a synthetic vinyl-type hydrophobic polymer, a non-vinyl-type hydrophobic polymer, naturally derived polymer, a membrane disruptive peptide, or a phospholipid bilayer disrupting agent.
 - 5-7. (Canceled)
- 8. (Currently amended) The composition of Claim [[33]] <u>36</u>, wherein the pH-sensitive linkage is an acetal, orthoester, cis-aconityl group, hydrazone, ester, Schiff base, vinyl ether, dithioacetal, tert butyl ester, carbamate, thioester, or phosphoramidate.
- 9. (Previously presented) The composition of Claim 34, wherein the therapeutic, diagnostic, or prophylactic agent is coupled to either the hydrophilic or the hydrophobic component by a degradable or disruptable linkage.
 - 10-12. (Canceled)
- 13. (Currently amended) The composition of Claim [[33]] <u>36</u>, wherein the conjugate further comprises a ligand, wherein the ligand specifically binds to a target molecule.
- 14. (Previously presented) The composition of Claim 34, wherein the therapeutic, diagnostic, or prophylactic agent is complexed to a component of the conjugate.
- 15. (Currently amended) The composition of Claim [[33]] <u>36</u>, wherein the pH sensitive linkage is hydrolyzed within about 30 to 60 minutes at a pH between 5.0 and 5.5.

- 16. (Currently amended) The composition of Claim [[33]] <u>36</u> further comprising a pharmaceutically acceptable carrier for delivery of the conjugate to a cell or organelle.
- 17. (Previously presented) The composition of Claim 16, wherein the carrier provides for systemic delivery of the conjugate, local delivery of the conjugate, or topical delivery of the conjugate.
 - 18. (Canceled)
- 19. (Previously presented) The composition of Claim 34, wherein the therapeutic, diagnostic, or prophylactic agent is an antisense nucleotide, ribozyme, ribozyme guide sequence, triplex forming oligonucleotide, or gene.

20-33. (Canceled)

- 34. (Currently amended) The composition of Claim [[33]] <u>36</u> further comprising an agent, wherein the agent is a therapeutic, diagnostic, or prophylactic agent.
- 35. (Currently amended) The composition of Claim [[33]] <u>36</u>, wherein the hydrophobic component comprises a synthetic polymer.
- 36. (Currently amended) [[The]] A composition of Claim 33, for enhancing transport through a membrane, comprising a hydrophilic conjugate having a hydrophobic component linked to a hydrophilic component by a pH-sensitive linkage, wherein the pH-sensitive linkage is stable at a pH between 6.8 and 8 and hydrolyzed at a pH less than 6.5 to release the hydrophobic component, wherein the hydrophilic component comprises a polyalkylene oxide, and wherein the hydrophobic component is membrane disruptive and allows enhanced transport through a membrane only when released from the hydrophilic conjugate.
 - 37. (Canceled)

38. (Previously presented) A conjugate, comprising:

(a) a hydrophobic synthetic vinyl-type polymer, wherein the polymer is

endosomal membrane disruptive when released from the hydrophilic conjugate:

(b) a plurality of pendant hydrophilic polyalkylene oxide components; and

(c) a plurality of pH-sensitive linkages, wherein each of the pendant

polyalkylene oxide components is covalently linked to the polymer through a pH-sensitive

linkage that is stable at a pH between 6.8 and 8 and hydrolyzed at a pH less than 6.5.

39. (Previously presented) The conjugate of Claim 38, wherein the synthetic

vinyl-type polymer is a terpolymer of dimethylaminoethyl methacrylate, butyl methacrylate, and

styrene benzaldehyde.

40. (Previously presented) The conjugate of Claim 38, wherein the pH-sensitive

linkage is selected from the group consisting of an acetal, a dithioacetal, an ester, an orthoester,

and a carbamate.

41. (Previously presented) A composition, comprising:

(a) a hydrophilic conjugate comprising:

(i) a hydrophobic synthetic vinyl-type polymer, wherein the polymer

is endosomal membrane disruptive when released from the hydrophilic conjugate;

(ii) a plurality of pendant hydrophilic polyalkylene oxide components;

and

(iii) a plurality of pH-sensitive linkages, wherein each of the pendant

polyalkylene oxide components is covalently linked to the polymer through a pH-sensitive

linkage that is stable at a pH between 6.8 and 8 and hydrolyzed at a pH less than 6.5; and

(b) a therapeutic or diagnostic agent.

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- 42. (Previously presented) The composition of Claim 41, wherein the synthetic vinyl-type polymer is a terpolymer of dimethylaminoethyl methacrylate, butyl methacrylate, and styrene benzaldehyde.
- 43. (Previously presented) The composition of Claim 41, wherein the pH-sensitive linkage is selected from the group consisting of an acetal, a dithioacetal, an ester, an orthoester, and a carbamate.
- 44. (Previously presented) The composition of Claim 41, wherein the therapeutic or diagnostic agent is selected from the group consisting of a protein, a peptide, a saccharide, a polysaccharide, an organic molecule, a nucleotide, an antisense nucleotide, an oligonucleotide, a ribozyme, a ribozyme guide sequence, a triplex forming oligonucleotide, and a gene.